

LORATADINE- loratadine tablet
Cabinet Health P.B.C.

Loratadine

Drug Facts

Active ingredient (in each tablet)

Loratadine USP 10 mg

Purpose

Antihistamine

Uses

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

• runny nose • itchy, watery eyes • sneezing • itching of the nose or throat

Warnings

Do not use

if you have ever had an allergic reaction to this product or any of its ingredients.

Ask a doctor before use if you have

liver or kidney disease. Your doctor should determine if you need a different dose.

When using this product

do not take more than directed. Taking more than directed may cause drowsiness.

Stop use and ask a doctor if

an allergic reaction to this product occurs. Seek medical help right away.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

adults and children 6 years and over	1 tablet daily; not more than 1 tablet in 24 hours
children under 6 years of age	ask a doctor
consumers with liver or kidney disease	ask a doctor

Other information

- Tamper-evident: do not use if foil seal under cap, printed with "SEALED for YOUR PROTECTION" is missing, open or broken
- store between 20° to 25°C (68° to 77°F)

Inactive ingredients

corn starch, lactose monohydrate, magnesium stearate, povidone

Questions or Comments?

Call + 1-908-242-6108

Package Labeling: 82725-0142-1

NDC 82725-0142-1
SKU CHB-10042-02A
Original Prescription Strength

Non-Drowsy*

Loratadine Tablets, USP

10 mg

Antihistamine **24 Hour Relief of:**

- Sneezing
- Runny Nose
- Itchy, Watery Eyes
- Itchy Throat or Nose

Indoor & Outdoor Allergies

***When taken as directed.
See Drug Facts Panel.**

8,000 Tablets

Drug Facts							
Active ingredient (in each tablet) Loratadine USP 10 mg	Purpose Antihistamine						
Uses temporarily relieves these symptoms due to hay fever or other upper respiratory allergies: ■ runny nose ■ itchy, watery eyes ■ sneezing ■ itching of the nose or throat							
Warnings Do not use if you have ever had an allergic reaction to this product or any of its ingredients. Ask a doctor before use if you have liver or kidney disease. Your doctor should determine if you need a different dose. When using this product do not take more than directed. Taking more than directed may cause drowsiness. Stop use and ask a doctor if an allergic reaction to this product occurs. Seek medical help right away. If pregnant or breast-feeding, ask a health professional before use. Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.							
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Questions or Comments? Call + 1-908-242-6108							

For Repackaging Only

Mfg. Lic. No. G/1430

1 4 1 7 2 1

Manufactured by:
Unique Pharmaceutical Laboratories
(A Div. of J. B. Chemicals & Pharmaceuticals Ltd.)
Mumbai 400 030, India

Manufactured for:
Cabinet Health P.B.C.
Brooklyn, NY 11222

NDC 82725-0142-3
SKU CHB-10042-02C
Original Prescription Strength

Non-Drowsy*

Loratadine Tablets, USP

10 mg

Antihistamine

Indoor & Outdoor Allergies

24 Hour Relief of:

• Sneezing

• Runny Nose

• Itchy, Watery Eyes

• Itchy Throat or Nose

*When taken as directed.

See Drug Facts Panel.

10,000 Tablets

142291

Drug Facts

Active ingredient (in each tablet)
Loratadine USP 10 mg

Purpose
Antihistamine

Uses

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:
■ runny nose ■ itchy, watery eyes ■ sneezing ■ itching of the nose or throat

Warnings

Do not use if you have ever had an allergic reaction to this product or any of its ingredients.

Ask a doctor before use if you have liver or kidney disease. Your doctor should determine if you need a different dose.

When using this product do not take more than directed. Taking more than directed may cause drowsiness.

Stop use and ask a doctor if an allergic reaction to this product occurs. Seek medical help right away.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

adults and children 6 years and over

1 tablet daily; not more than 1 tablet in 24 hours

children under 6 years of age

ask a doctor

consumers with liver or kidney disease

ask a doctor

Other information

■ store between 20° to 25°C (68° to 77°F)

Inactive ingredients

corn starch, lactose monohydrate, magnesium stearate, povidone

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Mfg. Lic. No. G/1430

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Manufactured for:

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Brooklyn, NY 11222

8 2 7 2 5 1 0 1 4 2 3

5

Package Labeling: 82725-0142-4

NDC 82725-0142-4
SKU CHB-10042-02D
Original Prescription Strength

Non-Drowsy*

Loratadine Tablets, USP

10 mg

Antihistamine 24 Hour Relief of:

- Sneezing
- Runny Nose
- Itchy, Watery Eyes
- Itchy Throat or Nose

Indoor & Outdoor Allergies

*When taken as directed.
See Drug Facts Panel.

Drug Facts	
Active ingredient (in each tablet)	Purpose
Loratadine USP 10 mg	Antihistamine
Uses	
temporarily relieves these symptoms due to hay fever or other upper respiratory allergies: ■ runny nose ■ itchy, watery eyes ■ sneezing ■ itching of the nose or throat	
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Other information	
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Questions or Comments?	
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For Repackaging Only

Mfg. Lic. No. G/1430

96,000 Tablets

142292



Manufactured by:
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Pharmaceuticals Ltd),
Mumbai 400 030, India

Manufactured for:
Cabinet Health P.B.C.
Brooklyn, NY 11222

Package Labeling: 82725-0142-2

NDC 82725-0142-2
SKU CHB-10042-02B
Original Prescription Strength

Non-Drowsy*

Loratadine Tablets, USP

10 mg

Antihistamine 24 Hour Relief of:

- Sneezing
- Runny Nose
- Itchy, Watery Eyes
- Itchy Throat or Nose

Indoor & Outdoor Allergies

*When taken as directed.
See Drug Facts Panel.

Drug Facts	
Active ingredient (in each tablet)	Purpose
Loratadine USP 10 mg	Antihistamine
Uses	
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Mfg. Lic. No. G/1430

100,000 Tablets 142292

LORATADINE

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82725-0142
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LORATADINE (UNII: 7AJ03BO7QN) (LORATADINE - UNII: 7AJ03BO7QN)	LORATADINE	10 mg

Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POVIDONE (UNII: FZ989GH94E)	

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	6mm
Flavor		Imprint Code	P;10
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82725-0142-1	8000 in 1 BAG; Type 0: Not a Combination Product	10/01/2025	
2	NDC:82725-0142-3	10000 in 1 BAG; Type 0: Not a Combination Product	10/01/2025	
3	NDC:82725-0142-4	1 in 1 DRUM	10/01/2025	
3		96000 in 1 DRUM; Type 0: Not a Combination Product		
4	NDC:82725-0142-2	1 in 1 DRUM	10/01/2025	
4		100000 in 1 BAG; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA214684	10/01/2025	

Labeler - Cabinet Health P.B.C. (117102391)

Revised: 10/2025

Cabinet Health P.B.C.